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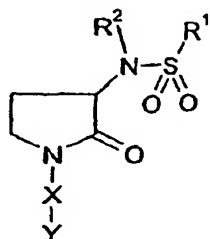
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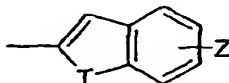
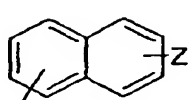
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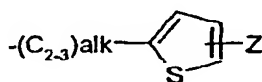
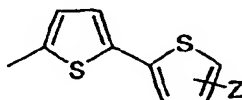
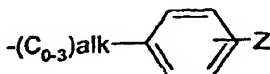
(54) Title: **1-PHENYL-2-OXO-3-SULFONYLAMINO-PYRROLIDINE DERIVATIVES AND RELATED COMPOUNDS AS FACTOR XA INHIBITORS FOR THE TREATMENT OF ACUTE VASCULAR DISEASES**



(I)



(II)



(57) Abstract: The invention relates to compounds of formula (I) wherein: R<sup>1</sup> represents a group selected from: formula (II) each ring of which optionally contains a further heteroatom N, Z represents an optional substituent halogen, alk represents alkylene or alkenylene, T represents S, O or NH; R<sup>2</sup> represents -C<sub>1-6</sub>alkyl, -C<sub>1-3</sub>alkylCN, -C<sub>0-3</sub>alkylR<sup>c</sup>, -C<sub>1-3</sub>alkylR<sup>f</sup>, -C<sub>2-3</sub>alkylNR<sup>a</sup>R<sup>b</sup>, -C<sub>2-3</sub>alkyl-OC<sub>1-6</sub>alkyl, -C<sub>2-3</sub>alkylOC<sub>1-3</sub>alkylCONR<sup>a</sup>R<sup>b</sup>, with the proviso that R<sup>2</sup> does not represent C<sub>2-3</sub>alkylmorpholino; X represents phenyl or a 5- or 6- membered aromatic heterocyclic group containing at least one heteroatom selected from O, N or S, each of which is optionally substituted by 0-2 groups selected from: halogen, -C<sub>1-4</sub>alkyl, -C<sub>2-4</sub>alkenyl, -CN, -CF<sub>3</sub>, -NR<sup>a</sup>R<sup>b</sup>, -C<sub>0-4</sub>alkylOR<sup>c</sup>, -C(O)R<sup>d</sup> and -C(O)NR<sup>a</sup>R<sup>b</sup>; Y represents a substituent selected from hydrogen, halogen, -C<sub>1-4</sub>alkyl, -C<sub>2-4</sub>alkenyl, -NR<sup>a</sup>R<sup>b</sup>, -NO<sub>2</sub>, -C(O)NR<sup>a</sup>R<sup>b</sup>, -N(C<sub>1-4</sub>alkyl)(CHO), -NHCOC<sub>1-4</sub>alkyl, -NHSO<sub>2</sub>R<sup>d</sup>, -C<sub>0-4</sub>alkylOR<sup>c</sup>, -C(O)R<sup>d</sup>, -S(O)<sub>n</sub>R<sup>d</sup>, or -S(O)<sub>2</sub>NR<sup>a</sup>R<sup>b</sup>; The other substituents are as defined in claim 1.

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